

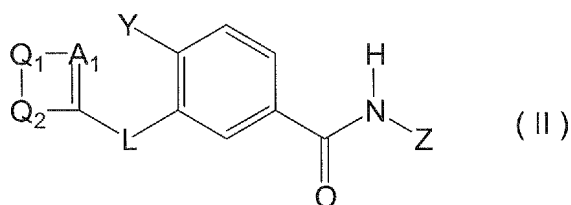
Amendments to the Claims

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

1. (Currently Amended) A compound of formula (II),
~~or a prodrug thereof, or a pharmaceutically acceptable salt~~
~~thereof of the compound or the prodrug.~~

~~{Formula 1}~~



where A₁ is C-X₁ ~~or N~~;

Q₁ is -A₂=A₃-~~, or a heteroatom selected from O, S,~~
~~, and N(R₁₀).~~;

Q₂ is -A₄=A₅-~~, or a heteroatom selected from O, S,~~
~~, and N(R₁₀).~~; provided that Q₁ and Q₂ are not heteroatoms at
the same time;

A₂ is C-X₂ ~~or N~~, A₃ is C-X₃ ~~or N~~, A₄ is C-X₄ ~~or N~~, and
A₅ is C-X₅ ~~or N~~;

R₁₀ is a hydrogen atom, C₁₋₆alkyl, haloC₁₋₆alkyl,
C₁₋₆alkylcarbonyl or aryl, the aryl being optionally

~~substituted by one or more substituents selected from a
halogen atom, C₁₋₆alkyl, and C₁₋₆alkoxy;~~

X₁, X₂, X₃, X₄ and X₅ are each independently selected from the group consisting of a hydrogen atom, hydroxy, a halogen atom, cyano, hydroxyaminocarbonyl, hydroxyamidino, nitro, amino, amidino, guanidino, C₁₋₆alkylamino, diC₁₋₆alkylamino, C₁₋₆alkylamidino, diC₁₋₆alkylamidino, C₁₋₆alkylguanidino, diC₁₋₆alkylguanidino, C₁₋₆alkylthio, C₁₋₆alkylsulfo, C₁₋₆alkylsulfonyl, C₁₋₆alkylphosphono, diC₁₋₆alkylphosphono, C₁₋₆alkyl, C₁₋₆alkoxy, C₃₋₉cycloalkyl, C₃₋₉cycloalkoxy, C₂₋₇alkenyl, C₂₋₇alkynyl, C₁₋₆alkylcarbonyl, C₁₋₆alkoxycarbonyl (the above 19 groups may be substituted by one or more substituents selected from a halogen atom, hydroxy, aryl, heteroaryl, and cyano), aryl, aryloxy, arylcarbonyl, heteroaryl, heteroaryloxy, heteroarylcarbonyl, and arylC₁₋₆alkyloxy (the above 7 groups may be substituted by one or more substituents selected from a halogen atom, C₁₋₆alkyl, and C₁₋₆alkoxy); or

X₁ and X₂, X₂ and X₃, X₃ and X₄, and X₄ and X₅, together with the carbon atoms to which they are bound, form a saturated or unsaturated 5- to 7-membered carbocyclic ring, or a saturated or unsaturated 5- to 7-membered heterocyclic ring containing one or more heteroatoms selected from an oxygen atom, a nitrogen atom, and a sulfur atom;

Y is selected from the group consisting of C₁₋₆alkyl, C₃₋₉cycloalkyl, C₂₋₇alkenyl, C₂₋₇alkynyl, C₁₋₆alkylcarbonyl, C₁₋₆alkoxycarbonyl, arylcarbonyl, heteroarylcarbonyl, aryloxy carbonyl, heteroaryloxy carbonyl, C₁₋₆alkoxy, C₂₋₇alkenyloxy, C₂₋₇alkynyloxy, C₁₋₆alkylthio, C₁₋₆alkylsulfonyl {the above 15 groups may be substituted by one or more substituents selected from a saturated or unsaturated 3- to 7-membered carbocyclyl, a saturated or unsaturated 3- to 7-membered heterocyclyl containing one or more heteroatoms selected from an oxygen atom, a nitrogen atom, and a sulfur atom, a halogen atom, hydroxy, C₁₋₆alkoxy, hydroxyC₁₋₆alkoxy, C₁₋₆alkoxyC₁₋₆alkoxy, aminoC₁₋₆alkoxy, N-C₁₋₆alkylaminoC₁₋₆alkoxy, N,N-diC₁₋₆alkylaminoC₁₋₆alkoxy, amino, C₁₋₆alkylamino, hydroxyC₁₋₆alkylamino, C₁₋₆alkoxyC₁₋₆alkylamino, aminoC₁₋₆alkylamino, diC₁₋₆alkylamino, bis(hydroxyC₁₋₆alkyl) amino, bis(C₁₋₆alkoxyC₁₋₆alkyl) amino, bis(aminoC₁₋₆alkyl) amino, amidino, C₁₋₆alkylamidino, diC₁₋₆alkylamidino, guanidino, C₁₋₆alkylguanidino, diC₁₋₆alkylguanidino, cyano, carboxyl, C₁₋₆alkoxycarbonyl, C₁₋₆alkylthio, C₁₋₆alkylsulfonyl, C₁₋₆alkylphosphono, and diC₁₋₆alkylphosphono}, amino, C₁₋₆alkylamino, diC₁₋₆alkylamino (the above 2 groups may be substituted by one or more substituents selected from a saturated or unsaturated 3- to 7-membered carbocyclyl, a saturated or unsaturated 3- to 7-membered heterocyclyl containing one or more heteroatoms

selected from an oxygen atom, a nitrogen atom, and a sulfur atom, a halogen atom, hydroxy, C₁₋₆alkoxy, hydroxyC₁₋₆alkoxy, C₁₋₆alkoxyC₁₋₆alkoxy, aminoC₁₋₆alkoxy, N-C₁₋₆alkylaminoC₁₋₆alkoxy, N,N-diC₁₋₆alkylaminoC₁₋₆alkoxy, amino, C₁₋₆alkylamino, hydroxyC₁₋₆alkylamino, C₁₋₆alkoxyC₁₋₆alkylamino, aminoC₁₋₆alkylamino, diC₁₋₆alkylamino, bis(hydroxyC₁₋₆alkyl)amino, bis(C₁₋₆alkoxyC₁₋₆alkyl)amino, bis(aminoC₁₋₆alkyl)amino, amidino, C₁₋₆alkylamidino, diC₁₋₆alkylamidino, guanidino, C₁₋₆alkylguanidino, diC₁₋₆alkylguanidino, cyano, carboxyl, C₁₋₆alkoxycarbonyl, C₁₋₆alkylthio, C₁₋₆alkylsulfonyl, C₁₋₆alkylphosphono, and diC₁₋₆alkylphosphono), a halogen atom, nitro, cyano, carboxyl, and a saturated or unsaturated 3- to 7-membered heterocyclyl containing one or more heteroatoms selected from an oxygen atom, a nitrogen atom, and a sulfur atom (the heterocyclyl may be substituted by one or more substituents selected from hydroxy, C₁₋₆alkyl, haloC₁₋₆alkyl, hydroxyC₁₋₆alkyl, C₁₋₆alkoxyC₁₋₆alkyl, and oxo);

Z is selected from the group consisting of a hydrogen atom, hydroxy, C₁₋₆alkyl, C₃₋₉cycloalkyl {the above 2 groups may be substituted by one or more substituents selected from a saturated or unsaturated 3- to 7-membered carbocyclyl (the carbocyclyl group may be substituted by one or more substituents selected from C₁₋₆alkyl, hydroxyC₁₋₆alkyl, and

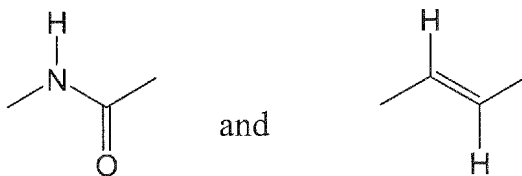
C₁₋₆alkoxyC₁₋₆alkyl), a saturated or unsaturated 3- to 7-membered heterocyclyl containing one or more heteroatoms selected from an oxygen atom, a nitrogen atom, and a sulfur atom (the heterocyclyl group may be substituted by one or more substituents selected from C₁₋₆alkyl, hydroxyC₁₋₆alkyl, and C₁₋₆alkoxyC₁₋₆alkyl), a halogen atom, hydroxy, C₁₋₆alkoxy, hydroxyC₁₋₆alkoxy, C₁₋₆alkoxyC₁₋₆alkoxy, hydroxyC₁₋₆alkoxyC₁₋₆alkoxy, aminoC₁₋₆alkoxy, N-C₁₋₆alkylaminoC₁₋₆alkoxy, N,N-diC₁₋₆alkylaminoC₁₋₆alkoxy, amino, C₁₋₆alkylamino, hydroxyC₁₋₆alkylamino, C₁₋₆alkoxyC₁₋₆alkylamino, aminoC₁₋₆alkylamino, diC₁₋₆alkylamino, bis(hydroxyC₁₋₆alkyl)amino, bis(C₁₋₆alkoxyC₁₋₆alkyl)amino, bis(aminoC₁₋₆alkyl)amino, cyano, carboxyl, C₁₋₆alkoxycarbonyl, aryloxy carbonyl, carbamoyl, C₁₋₆alkylcarbamoyl, diC₁₋₆alkylcarbamoyl{the above 2 groups may be substituted by one or more substituents selected from a halogen atom, hydroxy, cyano and amino), phosphono, C₁₋₆alkylphosphono, diC₁₋₆alkylphosphono, sulfonic acid, and C₁₋₆alkylsulfo}, and -OR₁ and -NR₁R₂;

R₁ and R₂ are each dependently selected from the group consisting of a hydrogen atom, C₁₋₆alkyl, C₁₋₆alkylcarbonyl, and a saturated or unsaturated 3- to 7-membered heterocyclyl containing one or more heteroatoms selected from an oxygen atom, a nitrogen atom, and a sulfur atom (the above 3 groups may be substituted by one or more

substituents selected from a saturated or unsaturated 3- to 7-membered carbocyclyl, a saturated or unsaturated 3- to 7-membered heterocyclyl containing one or more heteroatoms selected from an oxygen atom, a nitrogen atom, and a sulfur atom, a halogen atom, hydroxy, C₁₋₆alkoxy, hydroxyC₁₋₆alkoxy, C₁₋₆alkoxyC₁₋₆alkoxy, aminoC₁₋₆alkoxy, N-C₁₋₆alkylaminoC₁₋₆alkoxy, N,N-diC₁₋₆alkylaminoC₁₋₆alkoxy, amino, C₁₋₆alkylamino, hydroxyC₁₋₆alkylamino, C₁₋₆alkoxyC₁₋₆alkylamino, aminoC₁₋₆alkylamino, diC₁₋₆alkylamino, bis(hydroxyC₁₋₆alkyl)amino, bis(C₁₋₆alkoxyC₁₋₆alkyl)amino, bis(aminoC₁₋₆alkyl)amino, cyano, carboxyl, C₁₋₆alkoxycarbonyl, aryloxy carbonyl, phosphono, C₁₋₆alkylphosphono, diC₁₋₆alkylphosphono, sulfonic acid, and C₁₋₆alkylsulfo); or R₁ and R₂, together with the nitrogen atoms to which they are bound, form a saturated or unsaturated 5- to 7-membered heterocyclic ring containing one nitrogen atom and optionally further containing one or more heteroatoms selected from an oxygen atom, a nitrogen atom, and a sulfur atom; and

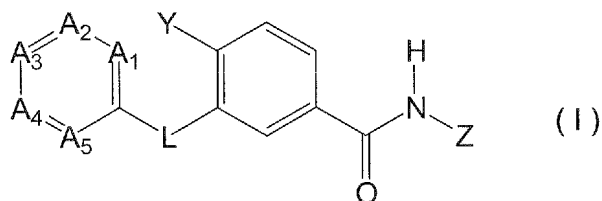
L is selected from the formula:

~~{Formula 2}~~



2. (Currently Amended) The compound, ~~or the prodrug thereof, or the pharmaceutically acceptable salt thereof of the compound or the prodrug,~~ according to claim 1, wherein the compound is represented by the formula (I):

~~{Formula 3}~~



where A_1 , A_2 , A_3 , A_4 , A_5 , L , Y , and Z are as defined in claim 1.

3. (Currently Amended) The compound, ~~or the prodrug thereof, or the pharmaceutically acceptable salt thereof of the compound or the prodrug,~~ according to claim 1 or 2, wherein Z is a hydrogen atom, C_{1-6} alkyl, C_{3-9} cycloalkyl, hydroxy C_{1-6} alkyl, hydroxy C_{1-6} alkoxy C_{1-6} alkyl, C_{1-6} alkoxy C_{1-6} alkyl, cyano C_{1-6} alkyl, pyridyl C_{1-6} alkyl, dihydroxy C_{1-6} alkyl, trihydroxy C_{1-6} alkyl, morpholino C_{1-6} alkyl, (N,N-di C_{1-6} alkylamino) C_{1-6} alkyl, or (N,N-bis(hydroxy C_{1-6} alkyl)amino) C_{1-6} alkyl.

4. (Currently Amended) The compound, ~~or the prodrug thereof,~~ or the pharmaceutically acceptable salt thereof ~~of the compound or the prodrug,~~ according to claim 3, wherein Z is a hydrogen atom, methyl, ethyl, cyclopropyl, cyclopentyl, 2-hydroxyethyl, 2-(2-hydroxyethoxy)ethyl, 2-methoxyethyl, 2-cyanoethyl, 4-pyridylmethyl, 1-methoxybut-2-yl, 2,3-dihydroxyprop-1-yl, 1,3-dihydroxyprop-2-yl, 1,3-dihydroxy-2-hydroxymethylprop-2-yl, 2-morpholinoethyl, 1-hydroxyprop-2-yl, 1-hydroxy-3-methylbut-2-yl, 2-(N,N-dimethylamino)ethyl, 2-(N,N-bis(2-hydroxyethyl)amino)ethyl, 2,4-dihydroxybutyl, 2,3,4-trihydroxybutyl, 2,3,4,5-tetrahydroxypentyl, or 2,3,4,5,6-pentahydroxyhexyl.

5. (Currently Amended) The compound, ~~or the prodrug thereof,~~ or the pharmaceutically acceptable salt thereof ~~of the compound or the prodrug,~~ according to any one of claims claim 1 to 4, wherein Y is a halogen atom, cyano, C₁₋₆alkyl, ~~haloC₁₋₆alkyl,~~ C₂₋₇alkenyl, C₂₋₇alkynyl, C₁₋₆alkoxy, C₃₋₉cycloalkylC₁₋₆alkoxy, C₂₋₇alkynyloxy, or haloC₁₋₆alkoxy.

6. (Currently Amended) The compound, ~~or the prodrug thereof,~~ or the pharmaceutically acceptable salt thereof ~~of the compound or the prodrug,~~ according to claim 5, wherein Y is chloro, bromo, cyano, methyl, ~~trifluoromethyl,~~

~~ethyl, n-propyl, i-propyl, ethynyl, methoxy, trifluoromethoxy,~~
cyclopropylmethoxy, 2-butyne-1-yloxy, or 2-chloroethoxy.

7. (Currently Amended) The compound, ~~or the~~
~~prodrug thereof, or the pharmaceutically acceptable salt~~
~~thereof of the compound or the prodrug,~~ according to claim 1
~~or 2,~~ wherein

~~A₁ is C-X₁ or N, A₂ is C-X₂ or N, A₃ is C-X₃ or N, A₄~~
~~is C-X₄ or N, and A₅ is C-X₅ or N,~~

X₁, X₂, X₃, X₄ and X₅ are each independently selected
from a hydrogen atom, a halogen atom, C₁₋₆alkyl, C₁₋₆alkoxy,
haloC₁₋₆alkyl, haloC₁₋₆alkoxy, C₁₋₆alkylthio, and
haloC₁₋₆alkylthio; or

X₁ and X₂, X₂ and X₃, X₃ and X₄, and X₄ and X₅,
together with the carbon atoms to which they are bound, form a
cyclohexane ring, a cyclopentane ring, a benzene ring, a
pyridine ring, a pyrimidine ring, a 1,4-dioxane ring, a 1,3-
dioxolane ring, a pyrrole ring, an imidazole ring, a thiazole
ring, or a furan ring.

8. (Currently Amended) The compound, ~~or the~~
~~prodrug thereof,~~ or the pharmaceutically acceptable salt
~~thereof of the compound or the prodrug,~~ according to claim 7,
wherein

X_1 , X_2 , X_3 , X_4 and X_5 are each independently selected from a hydrogen atom, fluoro, chloro, bromo, methyl, ethyl, t-butyl, i-propyl, methoxy, i-propoxy, trifluoromethyl, trifluoromethoxy, methylthio, and trifluoromethylthio; or

X_1 and X_2 , together with the carbon atoms to which they are bound, form a cyclohexane ring;

X_1 and X_2 , together with the carbon atoms to which they are bound, form a pyridine ring;

X_2 and X_3 , together with the carbon atoms to which they are bound, form a 1,4-dioxane ring; or

X_2 and X_3 , together with the carbon atoms to which they are bound, form a cyclopentane ring.

9-11. (Cancelled).

12. (Currently Amended) A pharmaceutical composition containing the compound, ~~or the prodrug thereof,~~ or the pharmaceutically acceptable salt thereof ~~of the compound or the prodrug,~~ according to ~~any one of claims claim~~ 1 to 11, as an active ingredient..

13. (Withdrawn-Currently Amended) An angiogenesis inhibitor containing the compound, ~~or the prodrug thereof,~~ or the pharmaceutically acceptable salt thereof ~~of the compound~~

~~or the prodrug, according to any one of claims claim 1 to 11,~~
as an active ingredient.

14. (Withdrawn-Currently Amended) An agent for treatment and prevention of a disease involving angiogenesis, said agent containing the compound, ~~or the prodrug thereof, or the pharmaceutically acceptable salt thereof of the compound~~ ~~or the prodrug, according to any one of claims claim 1 to 11,~~ as an active ingredient.

15. (Withdrawn) The agent for treatment and prevention, according to claim 14, wherein said disease involving angiogenesis is a cancerous disease.

16. (Withdrawn) The agent for treatment and prevention, according to claim 15, wherein said cancerous disease is solid tumor.

17. (Withdrawn-Currently Amended) An agent for treatment and prevention of metastasis of solid tumor, said agent containing the compound, ~~or the prodrug thereof, or the pharmaceutically acceptable salt thereof of the compound~~ ~~or the prodrug, according to any one of claims claim 1 to 11,~~ as an active ingredient.